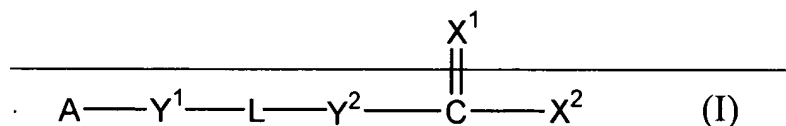


CLAIM AMENDMENTS

1. **(Currently Amended)** A method of inhibiting histone deacetylation activity in cells comprising contacting the cells with an effective amount of a compound wherein the compound is 5-phenyl-2,4-pentadienoic acid, 7-phenyl-2,4,6-heptatrienoic acid, 7-phenyl-2,4,6-heptatrienoylhydroxamic acid, 8-phenyl-3,5,7-octatrienoic acid, cinnamoylhydroxamic acid, methylcinnamoylhydroxamic acid, 5-phenyl-2,4-pentadienoylhydroxamic acid, N-methyl-5-phenyl-2,4-pentadienoylhydroxamic acid, 3-methyl-5-phenyl-2,4-pentadienoylhydroxamic acid, 4-methyl-5-phenyl-2,4-pentadienoyl hydroxamic acid, 4-chloro-5-phenyl-2,4-pentadienoylhydroxamic acid, 5-phenyl-2-en-4-yn-pentanoylhydroxamic acid, or N-methyl-6-phenyl-3,5-hexadienoylhydroxamic acid of formula (I), thereby treating one or more disorders mediated by histone deacetylase; said compound having the following formula:



wherein

~~A is a cyclic moiety selected from the group consisting of aryl, or heteroaryl; the cyclic moiety being optionally substituted with alkyl, alkenyl, alkynyl, alkoxy;~~

~~each of Y¹ and Y², independently, is a bond;~~

~~L is a straight C₂₋₁₂ hydrocarbon chain containing at least one double bond, at least one triple bond, or at least one double bond and one triple bond; said hydrocarbon chain being optionally substituted with C₁₋₄ alkyl, C₂₋₄ alkenyl, C₂₋₄ alkynyl, C₁₋₄ alkoxy, hydroxyl, halo, amino, nitro, cyano, C₃₋₅ cycloalkyl, 3-5 membered heterocycloalkyl, monocyclic aryl, 5-6 membered heteroaryl, C₁₋₄ alkylcarbonyloxy, C₁₋₄ alkylloxycarbonyl, C₁₋₄ alkylcarbonyl, or formyl; and further being optionally interrupted by O, N(R^e), N(R^e)-C(O)-O, -O-C(O)-N(R^e), N(R^e)-C(O)-N(R^f), or O-C(O)-O; each of R^e and R^f, independently, being hydrogen, alkyl, alkenyl, alkynyl, alkoxy, hydroxylalkyl, hydroxyl, or haloalkyl;~~

~~X¹ is O or S; and~~

~~X² is OR¹, SR¹, NR³OR¹, NR³SR¹, C(O)OR¹, CHR⁴OR¹, N=N-C(O)-N(R³)₂,
or O-CHR⁴-O-C(O)-R⁵, where each of R¹ and R², independently, is hydrogen, alkyl,
hydroxylalkyl, haloalkyl, or a hydroxyl protecting group; R³ is hydrogen, alkyl, alkenyl, alkynyl,
alkoxy, hydroxylalkyl, hydroxyl, haloalkyl, or an amino protecting group; R⁴ is hydrogen, alkyl,
hydroxylalkyl, or haloalkyl; and R⁵ is alkyl, hydroxylalkyl, or haloalkyl;
or a salt thereof; and~~

determining whether the level of acetylated histones in the treated cells is higher than in
untreated cells under the same conditions.

Claims 2-40. **(Cancelled)**

41. **(Currently Amended)** The method of claim 1, wherein said compound is ~~5-phenyl-2,4-~~
~~pentadienoic acid, 8-phenyl-3,5,7-octatrienoic acid, 5-phenyl-2,4-pentadienoylhydroxamic acid,~~
~~or 7-phenyl-2,4,6-hepta-trienoylhydroxamic acid.~~

Claims 42-43. **(Cancelled)**

44. **(Original)** The method of claim 1, wherein the cells being treated are cancerous.

45. **(Cancelled)**

46. **(Previously Presented)** The method of claim 1, wherein the disorder is cancer.

Claims 47-66 **(Cancelled)**

67. **(Currently Amended)** The method of claim [[40]] 1, wherein said compound is 5-
phenyl-2,4-pentadienoic acid.

Claims 68-73 **(Cancelled)**

74. **(Currently Amended)** The method of claim [[40]] 1, wherein said compound is 7-
phenyl-2,4,6-heptatrienoic acid.

75. **(Currently Amended)** The method of claim [[40]] 1, wherein said compound is 8-phenyl-3,5,7-octatrienoic acid.
76. **(Currently Amended)** The method of claim [[40]] 1, wherein said compound is cinnamoylhydroxamic acid.
77. **(Currently Amended)** The method of claim [[40]] 1, wherein said compound is methyl-cinnamoylhydroxamic acid.
78. **(Currently Amended)** The method of claim [[40]] 1, wherein said compound is 5-phenyl-2,4-pentadienoylhydroxamic acid.
79. **(Currently Amended)** The method of claim [[40]] 1, wherein said compound is N-methyl-5-phenyl-2,4-pentadienoylhydroxamic acid.
80. **(Currently Amended)** The method of claim [[40]] 1, wherein said compound is 3-methyl-5-phenyl-2,4-pentadienoylhydroxamic acid.
81. **(Currently Amended)** The method of claim [[40]] 1, wherein said compound is 4-methyl-5-phenyl-2,4-pentadienoyl hydroxamic acid.
82. **(Currently Amended)** The method of claim [[40]] 1, wherein said compound is 4-chloro-5-phenyl-2,4-pentadienoylhydroxamic acid.
83. **(Cancelled)**
84. **(Currently Amended)** The method of claim [[40]] 1, wherein said compound is 5-phenyl-2-en-4-yn-pentanoylhydroxamic acid.
85. **(Currently Amended)** The method of claim [[40]] 1, wherein said compound is N-methyl-6-phenyl-3,5-hexadienoylhydroxamic acid.